

(19)



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(11)

EP 1 388 531 A1

(12)

EUROPEAN PATENT APPLICATION(43) Date of publication:
11.02.2004 Bulletin 2004/07(51) Int Cl.7: **C07C 41/01**, C07C 41/06,
C07C 43/12(21) Application number: **03017180.5**(22) Date of filing: **29.07.2003**(84) Designated Contracting States:
AT BE BG CH CY CZ DE DK EE ES FI FR GB GR
HU IE IT LI LU MC NL PT RO SE SI SK TR
Designated Extension States:
AL LT LV MK

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20121 Milano (IT)(54) **Process for preparing fluorohalogenethers**

(57) A process for preparing (per) fluorohalogenethers having general formula (I):



wherein:

wherein:

A and A', equal to or different the one from the other, are Cl or Br or one is selected from A and A' and hydrogen and the other is halogen selected from Cl, Br; R = F, or a fluorinated, preferably perfluorinated, substituent, selected from the following groups: linear or branched C₁-C₂₀ alkyl more preferably C₁-C₁₀; C₃-C₇ cycloalkyl; aromatic, C₆-C₁₀ arylalkyl, alkylaryl; C₅-C₁₀ heterocyclic or alkylheterocyclic; when R is fluorinated or perfluorinated alkyl, cycloalkyl, arylalkyl, alkylaryl, it can optionally contain in the chain one or more oxygen atoms; when R is fluorinated it can optionally contain one or more H atoms and/or one or more halogen atoms different from F; n is an integer and is 1 or 2; m = 3-n;

p is an integer and is 1 or 2; q is an integer and is zero or 1; R is as above;

in liquid phase with elemental fluorine and with olefinic compounds having formula (III):



wherein A and A' are as above,
at temperatures in the range from -120°C to -20°C.

by reaction of carbonyl compounds having formula (II):

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Description

[0001] The present invention relates to a process for the preparation of fluorinated vinyl ethers.

[0002] More specifically the present invention refers to the fluorohalogen ether preparation which by dehalogenation produce the fluorinated vinyl ethers. The invention process leads to obtain fluorohalogen ethers having improved selectivities.

[0003] As known, fluorinated vinyl ethers are a class of valuable monomers to obtain various polymers, from fluorinated elastomers to thermoprocessable semicrystalline fluorinated polymers.

[0004] Processes to obtain fluorohalogen ethers based on the reaction of hypofluorites with olefins, are known in the prior art. For the hypofluorite preparation the most known processes use catalysts based on metal fluorides.

[0005] In USP 4,827,024 it is described the preparation in a continuous way of hypofluorite, by the fluorination reaction in equimolecular amounts with fluorine and halogenated carbonyl compounds having at least two carbon atoms, in the presence of catalysts formed of CsF as such or mixed with metals, such for example copper. Generally said metals are used, besides as catalyst (CsF) supports, also to make easier the thermal exchange, i.e. the dissipation of heat generated in the hypofluorite synthesis.

[0006] The metal support according to the above described prior art must satisfy two main functions: 1) to maintain the catalyst in a form accessible to reactants, 2) to make easier the thermal exchange maintaining under control in the required range the catalytic bed temperature. Last but not least a key feature of the support is the complete inertia towards reactants and reaction compounds.

[0007] In USP 4,816,599, USP 4,801,409 and USP 4,962,282 hypofluorites are preferably prepared with fluorine in excess to completely convert the acylfluoride into hypofluorite so that the acylfluoride concentration on the catalytic bed be very low, since it is known that some acylfluorides cause decomposition reactions in the presence of CsF. See for example Carl G. Ktrespan in Journal of Fluorine Chemistry, 16 (1980) 385-390.

[0008] Tests carried out by the Applicant on the prior art processes for the preparation of hypofluorites using the above catalysts have shown that by using both in a discontinuous and in a continuous way said catalytic systems, the catalytic activity rapidly decreases in the time. The Applicant has found in particular that the activity reduction is very marked, toll to the complete catalyst deactivation, when in the hypofluorite formation reaction the catalyst is used with an excess of fluorine over the stoichiometric value, the latter condition being indicated as preferred in the described prior art processes.

[0009] According to the prior art one must therefore operate in excess of fluorine in the hypofluorite synthesis to reduce as much as possible the above inconven-

iences. By operating under said conditions the catalyst of the prior art deactivates very rapidly, in two-three days. With so short durations it is in practice impossible to have available a continuous industrial plant.

[0010] Furthermore in discontinuous hypofluorite synthesis, when the catalytic bed is used in absence of support, its successive reuse in the hypofluorite obtainment reaction leads to very low yields and a very rapid deactivation is observed.

[0011] Processes to obtain fluorinated vinyl ethers are known in the prior art. USP 4,900,872 describes the perfluorovinylether precursor preparation, by continuous reaction between perfluoroalkyl hypofluorites diluted in an inert solvent and an olefin having formula $CA'F=CA''F$, wherein A and A', equal to or different from each other, are Cl and Br. In the patent it is indicated that said hypofluorites can be directly fed from the reactor wherein their synthesis in gaseous phase takes place, by reaction of fluorine with acylfluoride on catalyst. The obtained compounds are converted to perfluorovinylethers by dehalogenation with zinc. In said process the drawbacks are those reported above as to the hypofluorite preparation. In particular the drawback of said processes is due to the fact to have to synthesize and immediately use hypofluorites, which as known are unstable compounds, in particular when the number of carbon atoms of the hypofluorite perfluoroalkyl chain is higher than or equal to 2. Besides, in the hypofluorite synthesis it is known that one must use a catalyst, with the above drawbacks.

[0012] The need was therefore felt to have available a process for preparing fluorohalogen ethers overcoming the drawbacks of the prior art.

[0013] The Applicant has surprisingly and unexpectedly found that by using the process described herein after it is possible to solve said technical problem, and therefore to have available a continuous or discontinuous industrial process having a very high selectivity.

[0014] An object of the present invention is a process to prepare (per)fluorohalogen ethers having general formula (I):



wherein:

A and A', equal to or different the one from the other, are Cl or Br or one is selected from A and A' and hydrogen and the other is halogen selected from Cl, Br;

R can have the following meanings: F or a fluorinated, preferably perfluorinated, substituent, selected from the following groups: linear or branched C_1-C_{20} alkyl more preferably C_1-C_{10} ; C_3-C_7 cycloalkyl; aromatic, C_6-C_{10} arylalkyl, alkylaryl; C_5-C_{10} heterocyclic or alkylheterocyclic;

when R is fluorinated or perfluorinated alkyl, cycloalkyl, arylalkyl, alkylaryl it can optionally contain in the chain one or more oxygen atoms;

when R is fluorinated it can optionally contain one or more H atoms and/or one or more halogen atoms different from F;

n is an integer and is 1 or 2;

m is equal to 3-n;

by reaction of carbonyl compounds having formula (II):



wherein:

p is an integer and is 1 or 2;

q is an integer and is zero or 1.

with the proviso that when p = 2, q = 0; when p = 1,

q = 1;

R is as above;

in liquid phase with fluorine and with olefinic compounds having formula (III):



wherein A and A' are as above,

operating at temperatures from -120°C to -20°C, preferably from -100°C to -40°C, optionally in the presence of a solvent inert under the reaction conditions.

[0015] The fluorine used in the reaction can optionally be diluted with an inert gas such for example nitrogen or helium.

[0016] The formula (II) compounds which can be used are acylfluorides such for example COF_2 , CF_3COF , C_2F_5COF , C_3F_7COF , $C_7F_{15}COF$, $CF_3CF(OCF_3)CF_2CF_2COF$, $CF_3O(CF_2)_2COF$; ketones as hexafluoroacetone, perfluorodiisopropylketone, etc. Acylfluorides are preferred.

[0017] The formula (III) compounds are for example 1,2-dichloro-1,2-difluoroethylene (CFC1112), 1,2-dibromo-1,2-difluoroethylene, preferably CFC 1112.

[0018] The process according to the present invention is carried out in a single reactor and the reaction can be carried out in a semicontinuous or continuous way.

[0019] The semicontinuous process can be for example carried out by feeding gaseous fluorine in the reactor containing the formula (II) carbonyl compounds and the formula (III) olefinic compounds. The molar ratio (II)/(III) can range in a wide range, for example between 0.05 and 10. The fluorine feeding is continued until total olefin conversion. Said condition can be determined when the reaction exothermy is no longer noticed. In fact by carrying out the reaction of compounds (III) and (II) for example at -100°C, as soon as the reaction compounds

react with the elemental fluorine, there is exothermy and the temperature increases of about 5°-15°C. Therefore the reaction ends when for example compound (III) has been completely consumed. At this point the reactor temperature comes back to the initial temperature.

[0020] In the continuous process the gaseous fluorine and compounds (II), (III) are fed into the reactor, until reaching the steady state. In practice the reactants are fed into the reactor with established flow-rates and the reaction mixture is continuously drawn. The steady state is reached when the concentration of the three reactants and of the reaction compounds in the reactor is equal to the concentration of the reactants and reaction compounds outflowing from the reactor.

[0021] The molar ratios among the reactants are not particularly binding for the present invention process, for example the (II)/(III) molar ratio can range from 0.05 to 10 and F_2 /(III) between 0.05 and 10.

[0022] As solvents in the present invention process, compounds which are liquid and inert in the above temperature range can be used. Compounds selected from (per)fluorocarbons, (per)fluoroethers, (per)fluoropolyethers, perfluoroamines, or respective mixtures, can be for example used. The skilled man in the art is able to select from the above classes the compounds to be used as solvents on the basis of their physical properties.

[0023] The Applicant has surprisingly and unexpectedly found that the reaction among a formula (II) compound, the formula (III) olefin and elemental fluorine, in the above temperature range, directly produces formula (I) fluorohalogenethers having an improved selectivity. This result is quite surprising and unexpected. Besides in the invention process no catalyst is used, as on the contrary indicated in the prior art for obtaining fluorohalogenethers from the hypofluorite precursor. Therefore the catalyst absence in the invention process notably simplifies the process, particularly on an industrial scale. Indeed the costs of the catalyst regeneration, of its substitution and generally of the management of the plant section comprising the catalytic reactor are very onerous.

[0024] The Applicant has found that by using the process of the prior art to obtain fluorohalogenethers by reaction between the olefin and the hypofluorite obtained from the corresponding acylfluoride, a high amount of hypofluorite decomposes. The hypofluorite decomposition reaction increases with its molecular weight; in practice it takes place if the hypofluorite is different from the methyl hypofluorite. See the comparative Examples. With the invention process there is an improved selectivity even when fluorohalogenethers deriving from the formula (II) compounds are prepared, independently from the number of R carbon atoms. In the invention Examples when the compound (II) is acetylfluoride or propionylfluoride the selectivity in the fluorohalogenether is substantially of the same order. Besides, compared with the prior art processes which use a catalyst to form the

hypofluorites, the invention process has a higher productivity since the plant needs no stops for the catalyst regeneration or substitution. In the invention process the plant part concerning the catalyst preparation, the catalytic section running and the catalyst regeneration is therefore eliminated.

[0025] The Applicant has found that in the present invention process the decomposition products deriving from compound (II) are negligible. See the Examples.

[0026] Furthermore with the present invention process it is also possible to operate at compound (II) low conversions with high selectivity in the fluorohalogenether.

[0027] The formula (II) compounds, differently from hypofluorites which are unstable compounds, do not decompose in the reaction environment and can be recovered for example by distillation. When one operates with hypofluorites, this is not possible owing to the dangerousness of the use of said compounds both during the reaction and during the recovery. It is well known that when hypofluorites are used in the reaction they are let completely react without accumulation in the reaction environment.

[0028] The following Examples illustrate the invention with non limitative purposes.

EXAMPLE 1

Synthesis of $\text{CF}_3\text{-CF}_2\text{-CF}_2\text{OCFCI-CF}_2\text{Cl}$

[0029] 57 g of CFCI=CFCI (CFC 1112) and 15 g of $\text{CF}_3\text{-CF}_2\text{COF}$ (perfluoropropionyl fluoride, PFPF) are introduced in a 50 cc glass reactor equipped with mechanical stirrer and the solution is maintained at the temperature of -100°C .

[0030] By means of a bubbling inlet, fluorine is fed diluted with nitrogen (molar ratio fluorine/nitrogen 1 : 5) for 6.5 hours. During the fluorine feeding period a moderate reaction exothermy is noticed.

[0031] The gas outflowing from the reactor are let flow through a trap containing a fluorinated solvent and maintained at -80°C .

[0032] At the reaction end the solutions discharged from the reactor and from the trap are analyzed by gas chromatography.

[0033] The material reaction balance is 93.6%, calculated on the solution discharged from the reactor and on the compounds in the trap fluorinated liquid, with conversion of the olefin CFC 1112 and of PFPF of 100% and 61.25%, respectively.

[0034] The main PFPF reaction compounds are the following:

- A) $\text{CF}_3\text{-CF}_2\text{-CF}_2\text{OCFCI-CF}_2\text{Cl}$ (propyl adduct)
- B) $\text{CF}_3\text{-CF}_2\text{-CF}_2\text{O}(\text{CFCI})_3\text{-CF}_2\text{Cl}$

[0035] The selectivity is 73% for A and 17% for B) calculated with respect to PFPF.

[0036] Parallely to the main reaction there is also the fluorination reaction of CFC 1112 to $\text{CF}_2\text{Cl-CF}_2\text{Cl}$ (CFC 114) and fluorodimerization to $\text{CF}_2\text{Cl-CFCI}_3\text{-CF}_2\text{Cl}$ (CFC 1112 dimer); the selectivity with respect to CFC 1112 is 40% for each compound.

[0037] The olefin molar balance is about 98%.

[0038] The solution discharged from the reactor is distilled. The compound A) structure is confirmed by the ^{19}F -NMR analysis.

EXAMPLE 2 (comparative)

Synthesis of $\text{CF}_3\text{-CF}_2\text{-CF}_2\text{O-CFCI-CF}_2\text{Cl}$ according to the prior art

a) Hypofluorite $\text{CF}_3\text{-CF}_2\text{-CF}_2\text{OF}$ synthesis according to USP 4,827,024

[0039] 2.7 g NI/h of fluorine diluted with nitrogen (molar ratio fluorine/nitrogen 1/10) and 2.3 NI/h of $\text{CF}_3\text{-CF}_2\text{-COF}$ (PFPF) are fed into a 500 cc tubular metal reactor, filled with CsF catalyst, mixed with copper wires to disperse the reaction heat. The PFPF is converted in hypofluorite $\text{CF}_3\text{-CF}_2\text{-CF}_2\text{OF}$ with a yield of 99.5%.

b) Synthesis of $\text{CF}_3\text{-CF}_2\text{-CF}_2\text{O-CFCI-CF}_2\text{Cl}$ by reaction between the hypofluorite and CFC 1112 according to USP 4,900,872.

[0040] The hypofluorite produced in a) is fed in a CSTR type reactor (continuous stirred tank reactor), containing 121.2 g of CFCI=CFCI (CFC 1112) and 452 g of $\text{CF}_2\text{Cl-CF}_3$ (CFC 115) as reaction solvent, and maintained at the temperature of -90°C . The reaction is carried out for 4 hours and then the reactor is discharged and the solution is analyzed by gas chromatography.

[0041] The reaction mass balance is 84.3% and the selectivity in propyl adduct referred to the PFPF is 48.1%. The solution is distilled and the isolated compound confirms the analytical yield.

[0042] The amount by moles of the formed COF_2 , determined by acidimetric titration of the compound in the gaseous effluents (off gas), which gives an index of the hypofluorite decomposition by β -scission, is equal to 36% of the starting PFPF.

EXAMPLE 3 (comparative)

Synthesis of $\text{CF}_3\text{-CF}_2\text{O-CFCI-CF}_2\text{Cl}$ (ethyl adduct)

a) Hypofluorite synthesis

[0043] One proceeds as in step a) of the Example 2 (comparative), but feeding 2.3 NI/h $\text{CF}_3\text{-COF}$ (PFAF) to the catalytic reactor. The yield in hypofluorite $\text{CF}_3\text{-CF}_2\text{OF}$ is 99.6%.

b) Synthesis of $\text{CF}_3\text{-CF}_2\text{O-CFCI-CF}_2\text{Cl}$

[0044] One proceeds as in step b) of the Example 2 (comparative) but without using any solvent, by feeding into the CSTR reactor 300 g of CFC 1112. The reaction mass balance is 90%.

[0045] The solution discharged from the reactor is analyzed by gaschromatography and the yield in ethyl adduct $\text{CF}_3\text{-CF}_2\text{O-CFCI-CF}_2\text{Cl}$, calculated with respect to the fed hypofluorite, is 75%. The remaining 25% of the hypofluorite decomposes forming COF_2 .

[0046] The solution is distilled to recover the ethyl adduct. The recovered amount confirms the yield previously calculated by gaschromatography.

EXAMPLE 4Synthesis of $\text{CF}_3\text{-CF}_2\text{O-CFCI-CF}_2\text{Cl}$

[0047] 49 g of $\text{CF}_3\text{-COF}$ (PFAF) and 24.4 g of $\text{CF}_3\text{-CF}_2\text{O-CFCI-CF}_2\text{Cl}$ (CFC 1112) are introduced into the equipment described in the Example 1. The reaction mixture is maintained at the temperature of -87°C , and 1.5 NI/h of fluorine diluted with nitrogen (molar ratio fluorine/nitrogen 1/5) are fed therein for 129 minutes.

[0048] The reaction mass balance is 90.5% with a PFAF conversion of 10%. The gaschromatographic analysis shows that the main product is $\text{CF}_3\text{-CF}_2\text{O-CFCI-CF}_2\text{Cl}$ (ethyl adduct) and the selectivity referred to PFAF is 80%. The complement to 100 are by-products.

EXAMPLE 5Synthesis of $\text{CF}_3\text{-(CF}_2)_6\text{-CF}_2\text{O-CFCI-CF}_2\text{Cl}$

[0049] 19.65 g of $\text{CF}_3\text{-OCFCI-CF}_2\text{Cl}$ as solvent, 23.07 g of $\text{CF}_3\text{-CF}_2\text{O-CFCI-CF}_2\text{Cl}$ (CFC 1112) and 7.57 g of $\text{CF}_3\text{-(CF}_2)_6\text{-COF}$ are introduced into the equipment described in the Example 1. The reaction mixture is maintained at the temperature of -55°C and 1.5 NI/h of fluorine diluted with nitrogen (molar ratio fluorine/nitrogen 1/5) are fed therein for 110 minutes.

[0050] The reaction mass balance is 92.2%, the acylfluoride conversion 25%.

[0051] The discharged solution is analyzed by gaschromatography and the main product is the ether $\text{CF}_3\text{-(CF}_2)_6\text{-CF}_2\text{O-CFCI-CF}_2\text{Cl}$ which forms with a selectivity, referred to the starting acylfluoride, of 90%, the complement to 100 being by-products.

[0052] The solution is treated with water and the separated organic phase is distilled. The ^{19}F NMR and GC MS analyses confirm the compound structure as above indicated.

EXAMPLE 6Synthesis of $\text{CF}_3\text{CF(OCF}_3\text{)CF}_2\text{CF}_2\text{CF}_2\text{O-CFCI-CF}_2\text{Cl}$

[0053] The Example 1 is repeated introducing into the reactor 16.58 g of $\text{CF}_3\text{CF(OCF}_3\text{)CF}_2\text{CF}_2\text{COF}$ and 48.85 g of $\text{CF}_3\text{CF=CFCl}$ (CFC 1112). The reaction mixture is maintained at the temperature of -81°C and 1.5 NI/h of fluorine diluted with nitrogen (molar ratio fluorine/nitrogen 1/5) are fed therein. The test is carried out for 264 minutes and the material balance is 93.5% with a starting acylfluoride conversion equal to 41.8%. The discharged solution is analyzed by gaschromatography and the main compounds are the following perfluorohalogenethers:

- $\text{CF}_3\text{CF(OCF}_3\text{)CF}_2\text{CF}_2\text{CF}_2\text{O-CFCI-CF}_2\text{Cl}$,
- $\text{CF}_3\text{CF(OCF}_3\text{)CF}_2\text{CF}_2\text{CF}_2\text{O-(CFCI)}_3\text{-CF}_2\text{Cl}$,

which form with a selectivity, referred to the starting acylfluoride, of 85% and 7.4%, respectively.

[0054] The solution is treated with water and from the separated organic phase the ether $\text{CF}_3\text{CF}_2\text{CF(CCF}_3\text{)CF}_2\text{CF}_2\text{O-CFCI-CF}_2\text{Cl}$ is distilled whose structure is confirmed by ^{19}F NMR analysis.

[0055] The obtained compound subjected to dechlorination with zinc in dimethyl formamide, at 75°C , provides the corresponding vinyl ether $\text{CF}_3\text{CF}_2\text{CF(OCF}_3\text{)CF}_2\text{CF}_2\text{O-CF=CF}_2$.

EXAMPLE 7Synthesis of $\text{CF}_3\text{O-CFCI-CF}_2\text{Cl}$

[0056] Into the equipment described in the Example 1 maintained at the temperature of -100°C , are introduced 19.2 g of $\text{CF}_2\text{Cl-CF}_3$.

[0057] 1.5 NI/h of fluorine diluted with nitrogen (molar ratio fluorine/nitrogen 1/2.5), 1.5 NI/h of $\text{CF}_3\text{CF=CFCl}$ and 1.5 NI/h of COF_2 are then fed.

[0058] The reaction is continued for 2 hours and at the end the reactor is discharged. The reaction mixture is analyzed by gaschromatography.

[0059] It is found that the conversion of CFC 1112 is 69.0% and that of COF_2 57.0%.

[0060] The reaction mixture is distilled. 12.0 g of a compound having formula $\text{CF}_3\text{O-CFCI-CF}_2\text{Cl}$ (^{19}F NMR) are recovered.

[0061] Selectivity with respect to COF_2 is 66.7% and with respect to CFC 1112 is 44.6%.

[0062] It is found that together with the synthesis of the above compound, also the fluorination reaction of CFC 1112 to give CFC 114 and of fluorodimerization to $\text{CF}_2\text{Cl-CFCI-CFCI-CF}_2\text{Cl}$ took place in the reactor.

[0063] Selectivity calculated with respect to CFC 1112, for the product CFC 114 is 35.1% and for the CFC 1112 dimer is 7.3%.

[0064] The olefin molar balance is 99 %.

EXAMPLE 8

Synthesis of $(CF_3)_2-CF-O-CFCl-CF_2Cl$

[0065] Into the equipment of the Example 1, main- 5
tained at a temperature of $-80^\circ C$. 16.8 g of $CF_3C(O)CF_3$
are loaded. 1.7 Nl/h of fluorine diluted with nitrogen (mo-
lar ratio fluorine/nitrogen 1/2.5) and 1.5 Nl/h di CF -
 $Cl=CFCl$ are then fed.

[0066] The reaction is continued for 4 hours and at the 10
end the reactor is discharged. The reaction mixture is
analyzed by gaschromatography.

[0067] It is found that both the conversions of CFC
1112 and of $CF_3C(O)CF_3$ are 100%.

[0068] The reaction mixture is distilled. 19.5 g of a 15
compound having formula $(CF_3)_2-CF-O-CFCl-CF_2Cl$
(^{19}F NMR) are recovered.

[0069] Selectivity with respect to $CF_3C(O)CF_3$ is 57.0
% and with respect to CFC 1112 is of 20.9 %.

[0070] It is found that together with the synthesis of 20
the above compound, also the fluorination reaction of
CFC 1112 to give CFC 114 and of fluorodimerization to
 $CF_2Cl-CFCl-CFCl-CF_2Cl$ took place in the reactor.

[0071] Selectivity calculated with respect to CFC 25
1112, for the product CFC 114 is 62.7% and for the CFC
1112 dimer is 1.6 %.

[0072] The molar balance is 99 % for CFC 1112 and
90% for $CF_3C(O)CF_3$.

EXAMPLE 9

Synthesis of $CF_3-CF_2-CF_2-O-CHF-CF_2Cl$

[0073] Into the equipment of the Example 1, main- 35
tained at a temperature of $-90^\circ C$, 70.0 g of CF_3CF_2COF
and 11.3 g di $CHF=CFCl$ (CFC 1122a) are loaded. 1.5
Nl/h of fluorine diluted with nitrogen (molar ratio fluo-
rine/nitrogen 1/4) are then fed.

[0074] The reaction is continued for 1.5 hours and at 40
the end the reactor is discharged. The reaction mixture
is analyzed by gaschromatography.

[0075] It is found that the conversion of CFC 1122a is
100% and that of CF_3CF_2COF 20.5%.

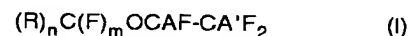
[0076] 1.2 g of $CF_3-CF_2-CF_2-O-CHF-CF_2Cl$ are re- 45
covered. Selectivity with respect to CF_3CF_2COF is 4.4
% and with respect to CFC 1122a is 3.4 %.

[0077] It is found that together with the synthesis of 50
the above compound, also the fluorination reaction of
CFC 1122a to give CFC 124a and of fluorodimerization
to $CF_2Cl-CFH-CFCl-CF_2H$ and isomers thereof
 $CF_2Cl-CFH-CFH-CF_2Cl$ and $CF_2H-CFCl-CFCl-CF_2H$.

[0078] Selectivity, calculated with respect to CFC 55
1122a, for CFC 124a is 39.4% and for CFC 1122a di-
mers is 13.6 %.

Claims

1. A process to prepare (per)fluorohalogenethers hav-
ing general formula (I):



wherein:

A and A', equal to or different the one from the
other, are Cl or Br or one is selected from A and
A' and hydrogen and the other is halogen sel-
ected from Cl, Br;

R = F or a fluorinated, preferably perfluorinated,
substituent, selected from the following groups:
linear or branched C_1-C_{20} alkyl more preferably
 C_1-C_{10} ; C_3-C_7 cycloalkyl; aromatic, C_6-C_{10} ar-
ylalkyl, alkylaryl; C_5-C_{10} heterocyclic or alkyl-
heterocyclic:

when R is fluorinated or perfluorinated
alkyl, cycloalkyl, arylalkyl, alkylaryl, it can
optionally contain in the chain one or more
oxygen atoms;

when R is fluorinated it can optionally con-
tain one or more H atoms and/or one or
more halogen atoms different from F;
n is an integer and is 1 or 2;

$$m = 3 - n;$$

by reaction of carbonyl compounds having formula
(II):



wherein:

p is an integer and is 1 or 2;

q is an integer and is zero or 1,

with the proviso that when $p = 2$, $q = 0$; when p
 $= 1$, $q = 1$;

R is as above;

in liquid phase with elemental fluorine and with
olefinic compounds having formula (III):



wherein A and A' are as above,
operating at temperatures from $-120^\circ C$ to $-20^\circ C$,
preferably from $-100^\circ C$ to $-40^\circ C$, optionally in the
presence of a solvent inert under the reaction con-
ditions.

2. A process according to claim 1, wherein the fluorine used in the reaction is diluted with an inert gas.
3. A process according to claims 1-2, wherein the formula (II) compounds are acylfluorides selected from COF_2 , CF_3COF , $\text{C}_2\text{F}_5\text{COF}$, $\text{C}_3\text{F}_7\text{COF}$, $\text{C}_7\text{F}_{15}\text{COF}$, $\text{CF}_3\text{CF}(\text{OCF}_3)$, $\text{CF}_2\text{CF}_2\text{COF}$, $\text{CF}_3\text{O}(\text{CF}_2)_2\text{COF}$; ketones selected between hexafluoroacetone, perfluorodiisopropylketone.
4. A process according to claims 1-4, wherein the formula (II) compounds are acylfluorides.
5. A process according to claims 1-4, wherein the formula (III) compounds are selected from 1,2-dichloro-1,2-difluoroethylene (CFC 1112), 1,2-dibromo-1,2-difluoro-ethylene, preferably CFC 1112.
6. A process according to claims 1-5, wherein the reaction can be carried out in a semicontinuous or continuous way.
7. A process according to claim 6, wherein in the semicontinuous process the molar ratio between the carbonyl compound (II) and the olefin (III) ranges from 0.05 to 10.
8. A process according to claim 6, wherein in the continuous process the molar ratio between the carbonyl compound (II) and the olefin (III) ranges from 0.05 to 10, the molar ratio fluorine/olefin (III) ranges from 0.05 to 10.

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EUROPEAN SEARCH REPORT

Application Number
EP 03 01 7180

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
D,Y	US 4 900 872 A (GUGLIELMO GIORGIO ET AL) 13 February 1990 (1990-02-13) * the whole document *	1-6	C07C41/01 C07C41/06 C07C43/12
D,Y	US 4 827 024 A (CARLINI FILIPPO M ET AL) 2 May 1989 (1989-05-02) * claim 1 *	1-6	
Y	EP 0 201 871 A (AUSIMONT SPA) 20 November 1986 (1986-11-20) * the whole document *	1-6	
A	US 3 453 333 A (LITT MORTON H ET AL) 1 July 1969 (1969-07-01) * claim 9; table 1 *	1	
A	PATENT ABSTRACTS OF JAPAN vol. 005, no. 085 (C-057), 3 June 1981 (1981-06-03) & JP 56 030935 A (ASAHI GLASS CO LTD), 28 March 1981 (1981-03-28) * abstract *	1	
			TECHNICAL FIELDS SEARCHED (Int.Cl.7)
			C07C
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 3 November 2003	Examiner Bonnevalle, E
CATEGORY OF CITED DOCUMENTS		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document	
X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document			

EPO FORM 1503 03.02 (P04C01)

**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

EP 03 01 7180

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

03-11-2003

Patent document cited in search report		Publication date		Patent family member(s)	Publication date
US 4900872	A	13-02-1990	IT	1198285 B	21-12-1988
			CA	1339538 C	11-11-1997
			DE	3767835 D1	07-03-1991
			EP	0267626 A1	18-05-1988
			JP	2512312 B2	03-07-1996
			JP	63198641 A	17-08-1988
US 4827024	A	02-05-1989	IT	1184149 B	22-10-1987
			AT	67176 T	15-09-1991
			AU	594326 B2	08-03-1990
			AU	5461586 A	16-10-1986
			CA	1271778 A1	17-07-1990
			DE	3681309 D1	17-10-1991
			EP	0194862 A2	17-09-1986
			IL	78104 A	05-11-1990
			JP	1927305 C	25-04-1995
			JP	6057679 B	03-08-1994
			JP	61257949 A	15-11-1986
			KR	9301326 B1	26-02-1993
			SU	1470176 A3	30-03-1989
			ZA	8601809 A	29-10-1986
EP 0201871	A	20-11-1986	IT	1187674 B	23-12-1987
			AT	36313 T	15-08-1988
			AU	584196 B2	18-05-1989
			AU	5737186 A	20-11-1986
			DE	3660490 D1	15-09-1988
			EP	0201871 A1	20-11-1986
			JP	62005932 A	12-01-1987
			SU	1533624 A3	30-12-1989
US 3453333	A	01-07-1969	BE	687649 A	01-03-1967
			DE	1568042 A1	26-03-1970
			FR	1500421 A	03-11-1967
			GB	1165912 A	01-10-1969
			GB	1165915 A	01-10-1969
			GB	1165916 A	01-10-1969
			NL	6613848 A	03-04-1967
			US	3542859 A	24-11-1970
JP 56030935	A	28-03-1981	US	3758538 A	11-09-1973
			NONE		

EPO FORM P0458

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82

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